

NYSTATIN- nystatin ointment

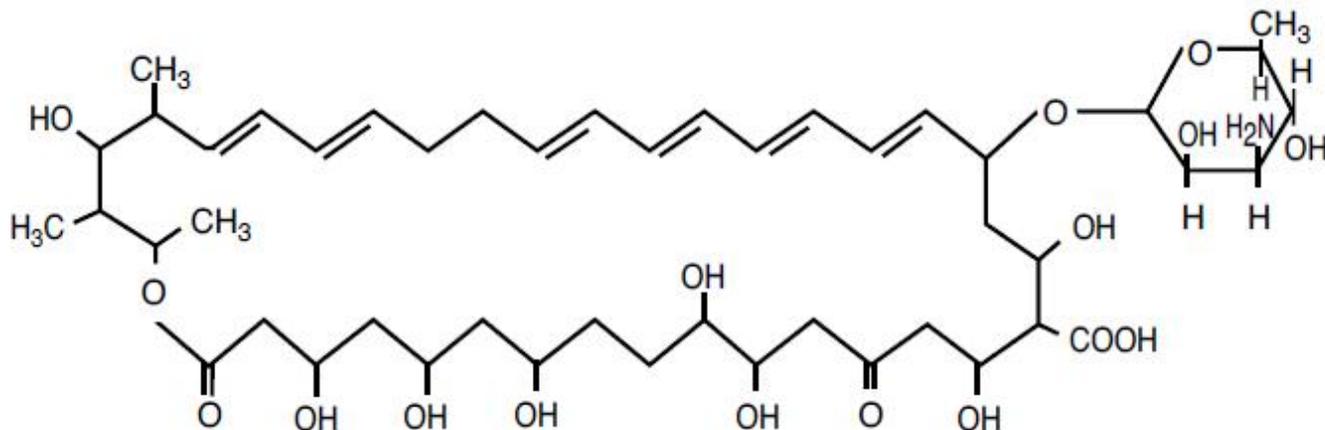
E. Fougera & Co. a division of Fougera Pharmaceuticals Inc.

NYSTATIN OINTMENT USP

R_x only

DESCRIPTION

Nystatin Ointment USP, for topical use only, contains 100,000 USP Nystatin Units per gram, in a polyethylene and mineral oil base. The structural formula is as follows:



Molecular Weight 926.13

Molecular Formula C₄₇H₇₅NO₁₇

CLINICAL PHARMACOLOGY

Nystatin is an antifungal antibiotic which is both fungistatic and fungicidal *in vitro* against a wide variety of yeasts and yeast-like fungi. It probably acts by binding to sterols in the cell membrane of the fungus with a resultant change in membrane permeability allowing leakage of intracellular components. Nystatin is a polyene antibiotic that is obtained from *Streptomyces noursei*, and is the first well tolerated antifungal antibiotic of dependable efficacy for the treatment of cutaneous, oral and intestinal infections caused by *Candida* [*Monilia*]*albicans* and other *Candida* species. It exhibits no appreciable activity against bacteria.

Nystatin provides specific therapy for all localized forms of candidiasis. Symptomatic relief is rapid, often occurring within 24 to 72 hours after the initiation of treatment. Cure is effected both clinically and mycologically in most cases of localized candidiasis.

INDICATIONS AND USAGE

Nystatin Ointment is indicated in the treatment of cutaneous or mucocutaneous mycotic infections caused by *Candida* [*Monilia*]*albicans* and other *Candida* species.

CONTRAINDICATIONS

Nystatin Ointment is contraindicated in patients with a history of hypersensitivity to any of its

components.

PRECAUTIONS

Should a reaction of hypersensitivity occur the drug should be immediately withdrawn and appropriate measures taken.

This preparation is not for ophthalmic use.

ADVERSE REACTIONS

Nystatin is virtually non-toxic and nonsensitizing and is well tolerated by all age groups including debilitated infants, even on prolonged administration. If irritation on topical application should occur, discontinue medication.

DOSAGE AND ADMINISTRATION

Nystatin Ointment should be applied liberally to affected areas twice a day or as indicated until healing is complete. Nystatin cream is usually preferred to nystatin ointment in candidiasis involving intertriginous areas; very moist lesions, however, are best treated with nystatin topical powder.

This preparation does not stain skin or mucous membranes and provides a simple, convenient means of treatment.

HOW SUPPLIED

Nystatin Ointment (100,000 USP Nystatin Units per gram) is a yellow ointment available as follows:

NDC 0168-0007-15 15 gram tube

NDC 0168-0007-30 30 gram tube

Store at controlled room temperature 15°- 30°C (59°-86°F).

E. FOUGERA & CO.

A division of Fougera Pharmaceuticals Inc.

Melville, New York 11747

I20715H

R08/11

#21

PACKAGE LABEL – PRINCIPAL DISPLAY PANEL – 15 G CONTAINER

NDC 0168-0007-15

FOUGERA®

NYSTATIN OINTMENT USP

100,000 USP Nystatin Units Per Gram

For Topical Use Only.

Not for Ophthalmic Use.

Rx only

CONTAINS: 100,000 USP

Nystatin Units per gram,

in a polyethylene and

mineral oil base.

NET WT 15 grams



PACKAGE LABEL – PRINCIPAL DISPLAY PANEL – 15 G CARTON

NDC 0168-0007-15

FOUGERA[®] Rx only

NYSTATIN OINTMENT USP

100,000 USP Nystatin Units Per Gram

**WARNING: Keep this and all
drugs out of reach of children.**

For Topical Use Only.

Not for Ophthalmic Use.

NET WT 15 grams



NYSTATIN

nystatin ointment

Product Information

| | | | |
|--------------------------------|-------------------------|---------------------------|---------------|
| Product Type | HUMAN PRESCRIPTION DRUG | Item Code (Source) | NDC:0168-0007 |
| Route of Administration | TOPICAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|-----------------------|
| nystatin (UNII: BDF1O1C72E) (nystatin - UNII:BDF1O1C72E) | nystatin | 100000 [USP'U] in 1 g |

Inactive Ingredients

| Ingredient Name | Strength |
|--|----------|
| HIGH DENSITY POLYETHYLENE (UNII: UG00KM4WR7) | |
| Mineral Oil (UNII: T5L8T28FGP) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:0168-0007-15 | 15 g in 1 TUBE; Type 0: Not a Combination Product | 09/23/1982 | |
| 2 | NDC:0168-0007-30 | 30 g in 1 TUBE; Type 0: Not a Combination Product | 09/23/1982 | |

Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| ANDA | ANDA062124 | 09/23/1982 | |

Labeler - E. Fougera & Co. a division of Fougera Pharmaceuticals Inc. (043838424)

Revised: 11/2018

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